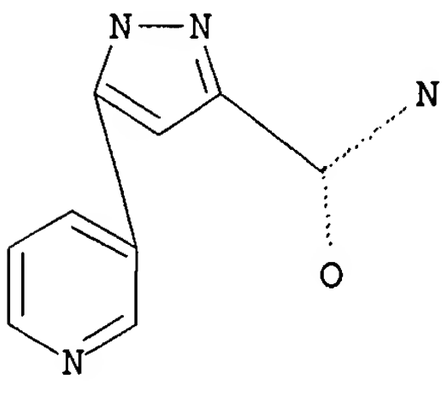
=> file caplus
FILE 'CAPLUS' ENTERED AT 12:50:06 ON 28 JUL 2004

FILE COVERS 1907 - 28 Jul 2004 VOL 141 ISS 5
FILE LAST UPDATED: 27 Jul 2004 (20040727/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d que
L1 STR



Structure attributes must be viewed using STN Express query preparation.
L3 17 SEA FILE=REGISTRY SSS FUL L1
L4 11 SEA FILE=CAPLUS L3

=> d l4 1-11 ibib abs hitstr

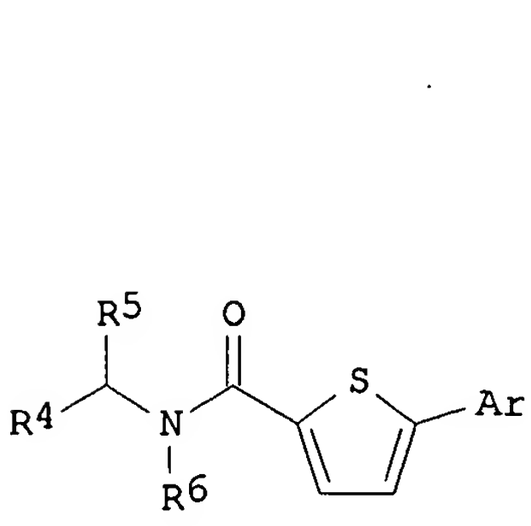
L4 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:493566 CAPLUS
DOCUMENT NUMBER: 141:38610
TITLE: Preparation of substituted thiophenes and related compounds as prenylation inhibitors
INVENTOR(S): Li, Francine Feirong; Rehder, Kenneth S.; Campbell, Michael Gordon; Viscardi, Celeste Patrice; Strachan, Jon-paul; Guo, Zhengming
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 117 pp., Cont.-in-part of U.S. Ser. No. 336,285.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 5
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004116425	A1	20040617	US 2003-636327	20030806
US 6649638	B1	20031118	US 2003-336285	20030103

PRIORITY APPLN. INFO.:

US 2002-219628 B2 20020814
US 2003-336285 A2 20030103
US 2003-454554P P 20030314

GI



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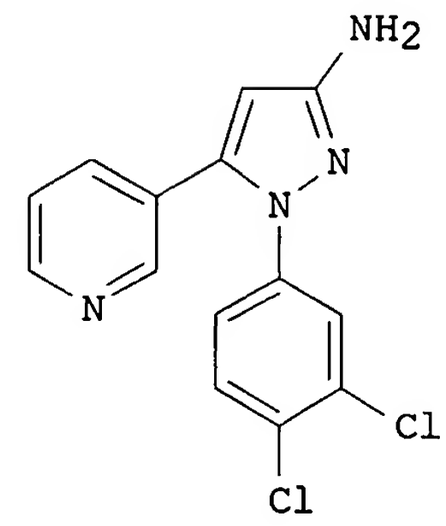


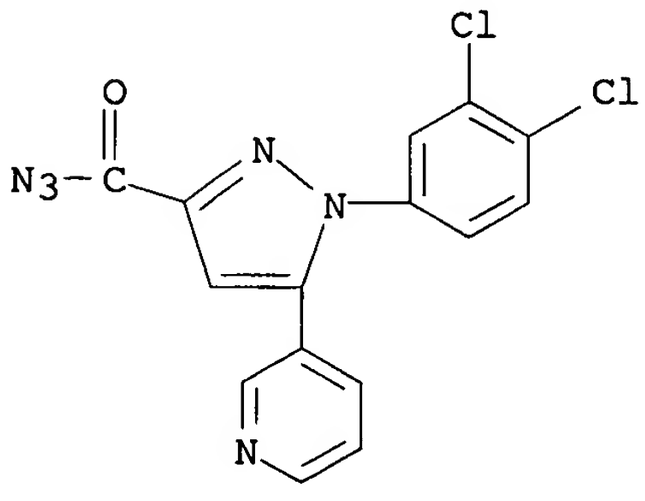
II

AB Title compds. I [Ar = heterocyclyl; R4 = absent, H, NH2, CONMe2, etc.; R5 = absent, i-Pr, benzyl, etc.; R6 = H, Me, Et, Pr, etc.] and related compds. are prepd. For instance, 1-(3,4-dichlorophenyl)-5-(pyridin-3-yl)-1H-pyrazole-3-carboxylic acid Me ester.bul.HCl (prepn. given) is sapond. (THF/H2O, NaOH) and converted to the Boc-protected pyrazole-3-amine (i. DMF, t-BuOH, DPPA, Et3N; ii. t-BuOH, reflux, 4 h) and deprotected to II. Compds. of the invention have inhibitory activity for GTPase I [no data]. I inhibit protein prenylation and are useful for treating cancer, restenosis, psoriasis, etc.

IT **623158-60-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of substituted thiophenes and related compds. as prenylation inhibitors)

RN 623158-60-1 CAPLUS

CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-(9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:162776 CAPLUS
DOCUMENT NUMBER: 140:217636
TITLE: Preparation of pyridinylpyrazolylcyclobutylacetamides

as prenylation inhibitors

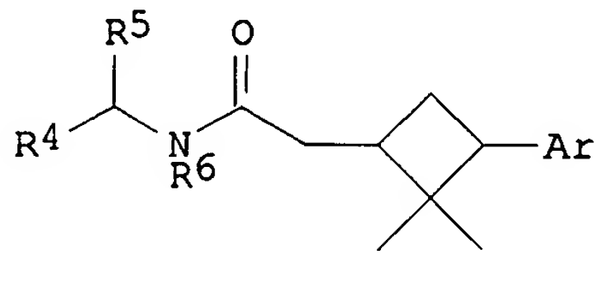
INVENTOR(S): Brown, Bradley B.; Rehder, Kenneth S.; Strachan, Jon-paul; Eaves, Jeron H.; Lowden, Christopher T.
PATENT ASSIGNEE(S): Ppd Discovery, Inc., USA
SOURCE: PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 5
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004016741	A2	20040226	WO 2003-US24984	20030806
WO 2004016741	A3	20040408		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 6664277	B1	20031216	US 2003-336186	20030103

PRIORITY APPLN. INFO.: US 2002-219851 A 20020814
US 2003-336186 A 20030103
US 2003-454554P P 20030314
OTHER SOURCE(S): MARPAT 140:217636
GI



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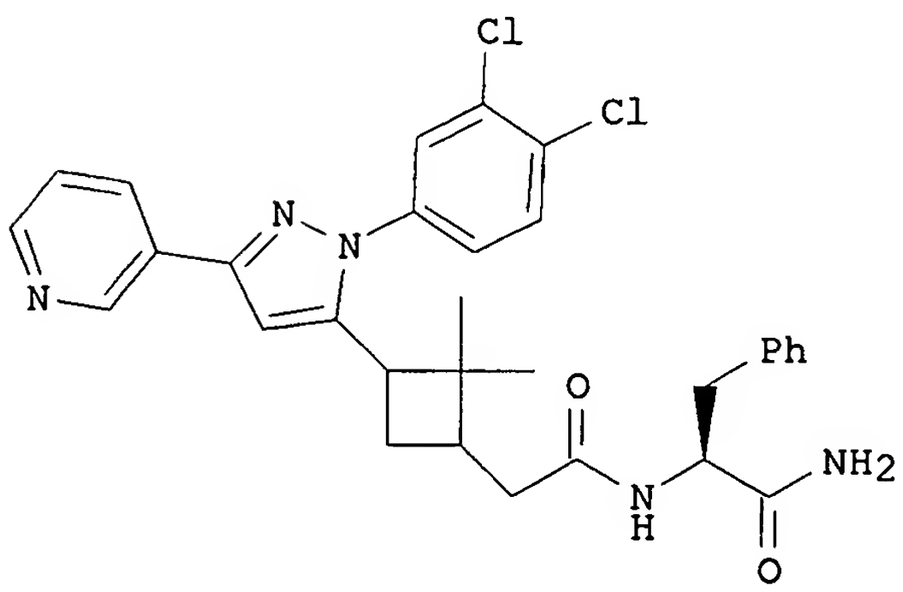


II

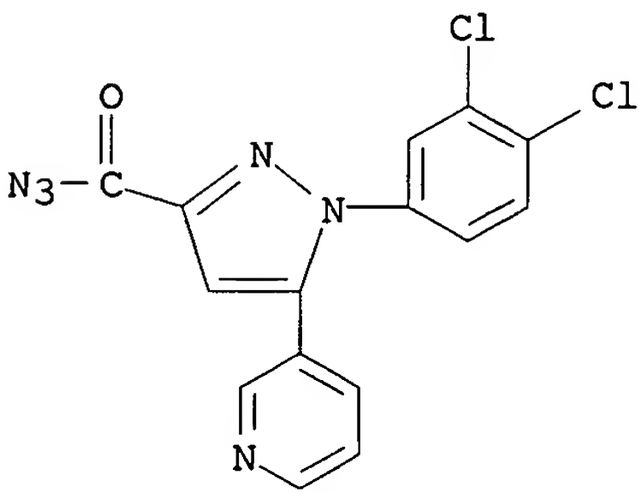
AB Title compds. [I; Ar = Q1, Q2; X = C, N, O, S; R1 = Ph, PhCH2, Me, Et, Pr, pyrimidinyl, etc.; R2 = Me, pyridyl, 3-cyanophenyl, 2-methylthiazolyl,

5-methylisoxazolyl, NMe2, etc.; R3 = null, CH2CH2OH, CH2CH2OMe, CH2CH2NMe2, CH2CH2CO2H, CH2OH, CH2CH2SMe, etc.; R4 = null, H, NH2, CONMe2, CO2H, cyano, CH2OH, CONHMe, CO2Me, C(:NOH)NH2, tetrazolyl, etc.; R5 = null, Me2CH, PhCH2, 4-trifluoromethylbenzyl, 4-cyanobenzyl, 3,4-dichlorobenzyl, 4-fluorobenzyl, etc.; R6 = H, Me, Et, Pr, Me2CH, CH2CO2H, PhCH2, CH2CO2Et, 2-methoxynaphthylmethyl], were prepd. for treatment of cancer, infection, ischemia, restenosis, psoriasis, endometriosis, atherosclerosis, hypercholesterolemia, angiogenesis, and corneal neovascularization (no data). Thus, title compd. (II) was prepd. in several steps from (S)-.alpha.-pinene.

IT **623158-60-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of pyridinylpyrazolylcyclobutylacetamides as prenylation inhibitors)

RN 623158-60-1 CAPLUS

CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:162671 CAPLUS
DOCUMENT NUMBER: 140:199323
TITLE: Preparation of substituted thiophenes and related compounds as prenylation inhibitors
INVENTOR(S): Li, Francine Feirong; Rehder, Kenneth S.; Campbell, Michael Gordon; Viscardi, Celeste Patrice; Strachan, Jon-Paul; Guo, Zhengming
PATENT ASSIGNEE(S): PPD Discovery, Inc., USA
SOURCE: PCT Int. Appl., 137 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 5
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004016592	A1	20040226	WO 2003-US24985	20030806

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,

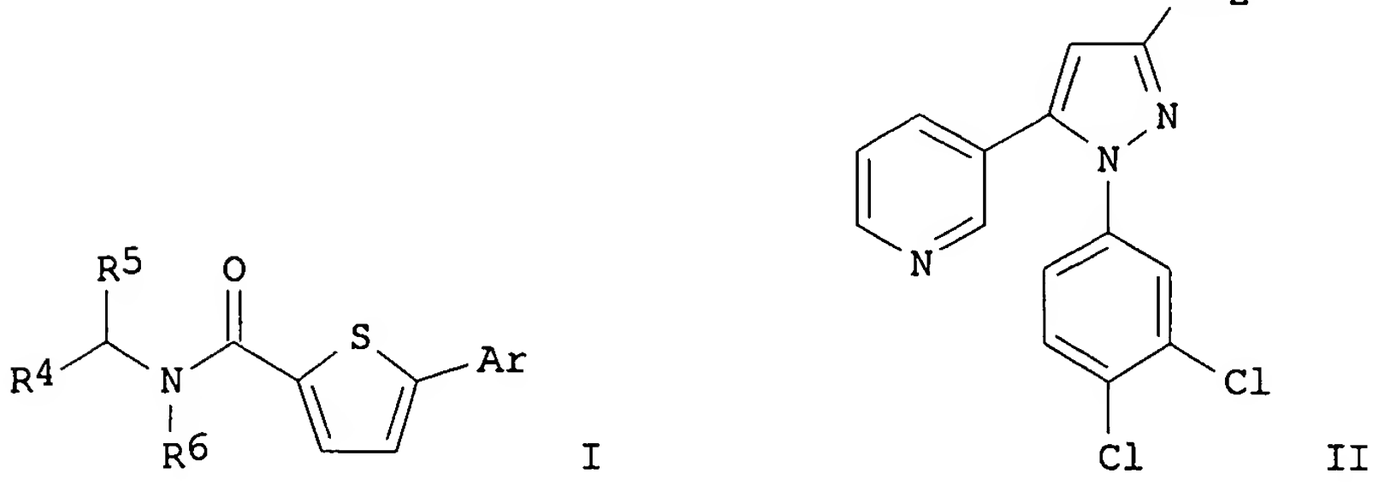
CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 6649638 B1 20031118 US 2003-336285 20030103

PRIORITY APPLN. INFO.: US 2002-219628 A 20020814
US 2003-336285 A 20030103
US 2003-454554P P 20030314

OTHER SOURCE(S): MARPAT 140:199323

GI



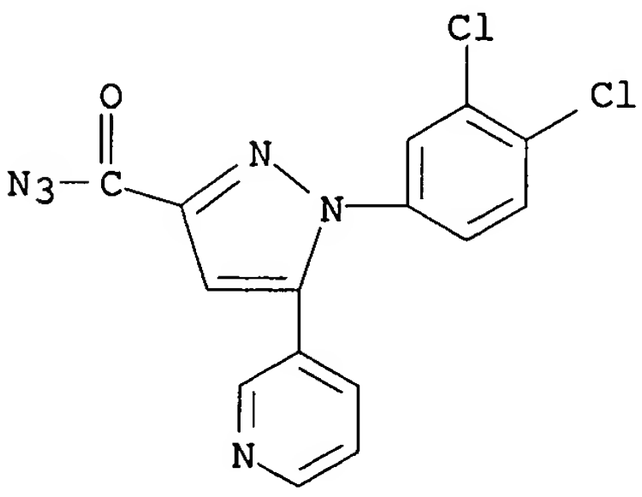
I II

AB Title compds. I [Ar = heterocyclyl; R4 = absent, H, NH2, CONMe2, etc.; R5 = absent, i-Pr, Benzyl, etc.; R6 = H, Me, Et, Pr, etc.] and related compds. are prepd. For instance, 1-(3,4-dichlorophenyl)-5-(pyridin-3-yl)-1H-pyrazole-3-carboxylic acid Me ester.bul.HCl (prepn. given) is sapond. (THF/H2O, NaOH) and converted to the Boc-protected pyrazole-3-amine (i. DMF, t-BuOH, DPPA, Et3N; ii. t-BuOH, reflux, 4 h) and deprotected to II. Compds. of the invention have inhibitory activity for GTPase I [no data]. I inhibit protein prenylation and are useful for treating cancer, restenosis, psoriasis, etc.

IT **623158-60-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of substituted thiophenes and related compds. as prenylation inhibitors)

RN 623158-60-1 CAPLUS

CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:930982 CAPLUS
DOCUMENT NUMBER: 140:782
TITLE: Methods using sulfonamide-containing cyclic compounds for treating carbonic anhydrase-mediated disorders
INVENTOR(S): Masferrer, Jaime L.; O'Neal, Janet M.
PATENT ASSIGNEE(S): Pharmacia Corporation, USA
SOURCE: U.S. Pat. Appl. Publ., 43 pp., Cont.-in-part of U.S. Ser. No. 213,793.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003220376	A1	20031127	US 2003-367384	20030214
US 2003100594	A1	20030529	US 2002-213793	20020807
WO 2004014430	A1	20040219	WO 2003-US4469	20030214

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

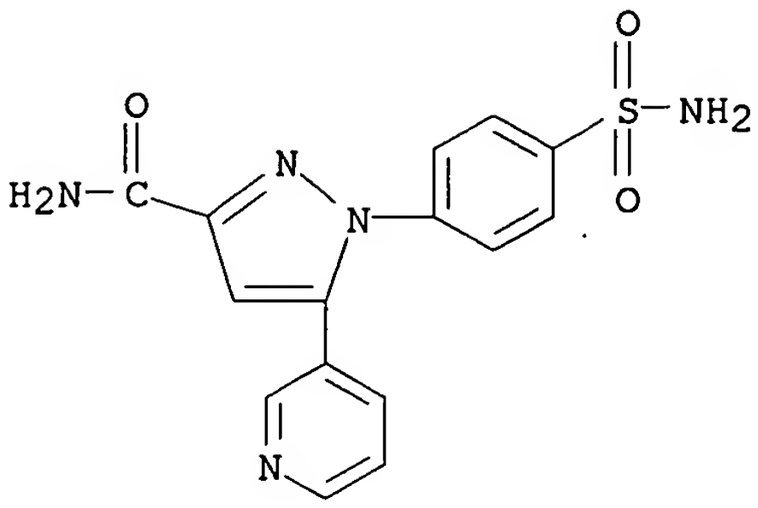
WO 2004014352	A2	20040219	WO 2003-US4494	20030214

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2001-311561P P 20010810
US 2002-213793 A2 20020807
OTHER SOURCE(S): MARPAT 140:782

AB The invention provides methods to treat or prevent carbonic anhydrase-mediated diseases or disorders. The method generally comprises administering a cyclic compd. having a sulfonamide group to a subject, wherein the compd. inhibits carbonic anhydrase.

IT **627094-49-9**
RL: AGR (Agricultural use); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(sulfonamide-contg. cyclic compds. for treating carbonic anhydrase-mediated disorders, and use with other agents)
RN 627094-49-9 CAPLUS
CN 1H-Pyrazole-3-carboxamide, 1-[4-(aminosulfonyl)phenyl]-5-(3-pyridinyl)-(9CI) (CA INDEX NAME)

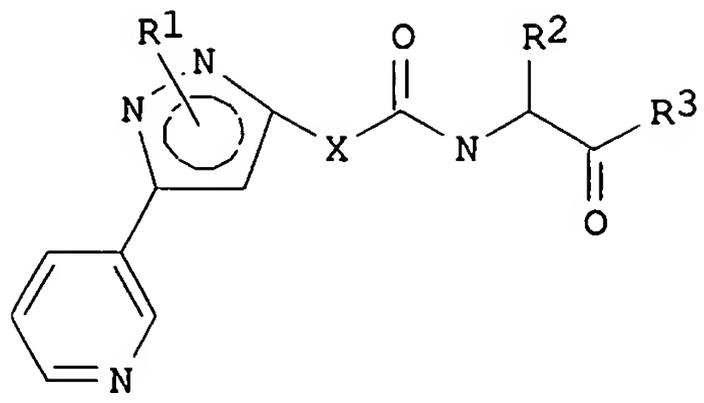


L4 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:903255 CAPLUS
DOCUMENT NUMBER: 139:396168
TITLE: Preparation of 3-pyridylpyrazole peptide derivatives as prenylation inhibitors
INVENTOR(S): Brown, Bradley B.; Rehder, Kenneth S.
PATENT ASSIGNEE(S): PPD Discovery, Inc., USA
SOURCE: U.S., 17 pp., Cont.-in-part of U.S. Ser. No. 219,628, abandoned.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 5
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6649638	B1	20031118	US 2003-336285	20030103
WO 2004016592	A1	20040226	WO 2003-US24985	20030806
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004116425	A1	20040617	US 2003-636327	20030806
US 2004053970	A1	20040318	US 2003-646256	20030822

PRIORITY APPLN. INFO.: US 2002-219628 B2 20020814
US 2003-336285 A 20030103
US 2003-454554P P 20030314

GI



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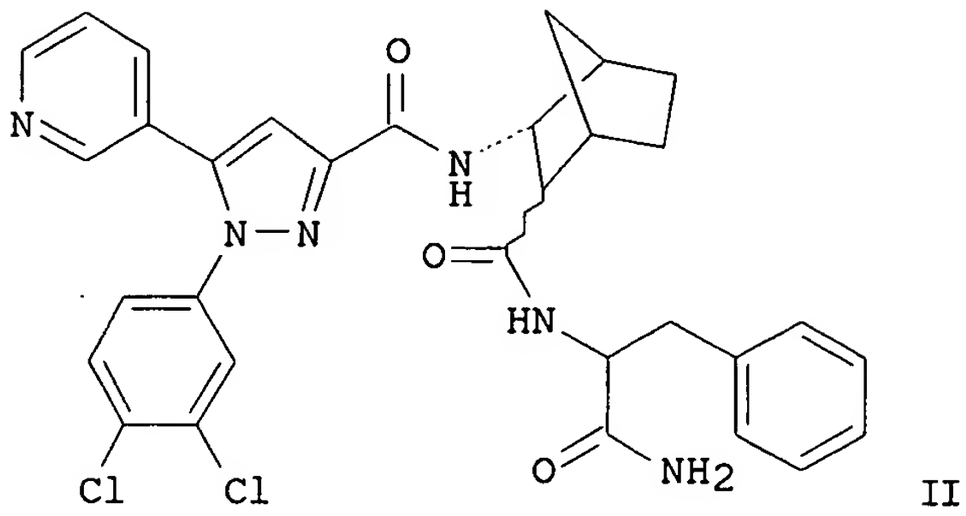


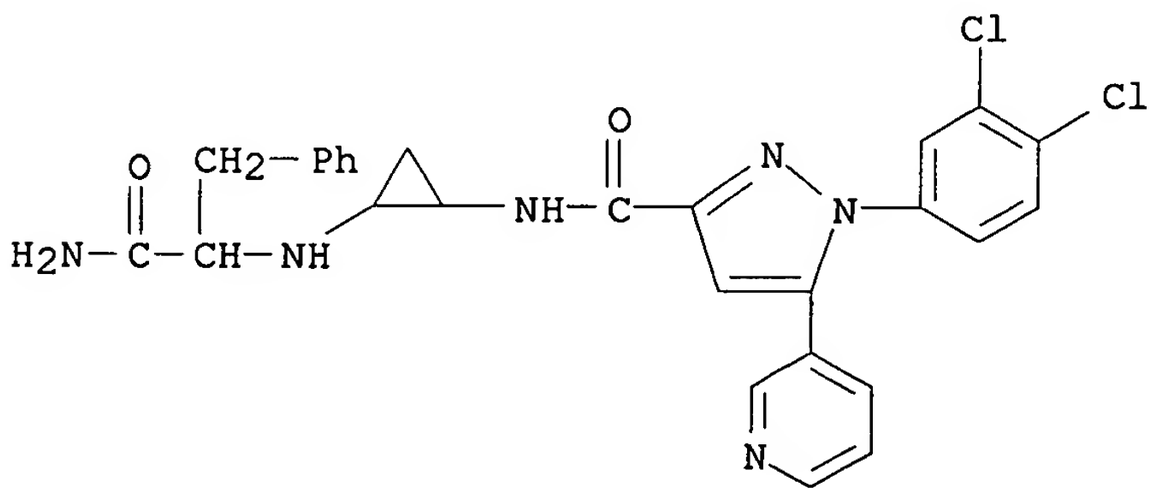
II

AB The invention is directed to pyridylpyrazole compds. I [X is nitrogen, Ph, pyrazole, methylpyrazole, dimethylpyrazole, pyridine, thiophene, dimethylcyclobutyl, dimethylcyclopropyl or cyclopropyl; R1 is halophenyl; R2 is benzyl, iso-Pr, chlorobenzyl, methylthienyl, (trifluoromethyl)benzyl, ethylthiomethyl, or 1-benzyl-4-pyrazolylmethyl; R3 is NH2 or OH] for use in the treatment of diseases assocd. with prenylation of proteins. Thus, phenylalaninamide deriv. II was prepd. via peptide coupling reactions and shown to inhibit GGPTase I.

IT **623158-71-4P 627088-86-2P 627088-99-7P**
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)

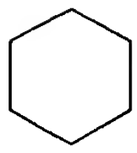
RN 623158-71-4 CAPLUS

CN 1H-Pyrazole-3-carboxamide, N-[2-[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]cyclopropyl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)

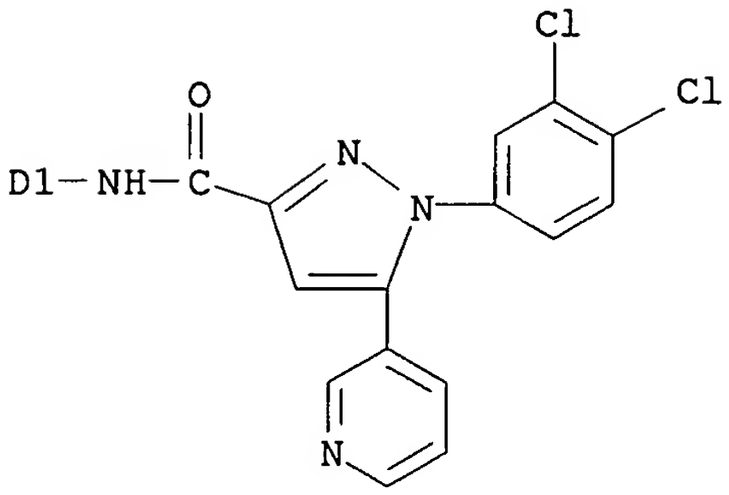


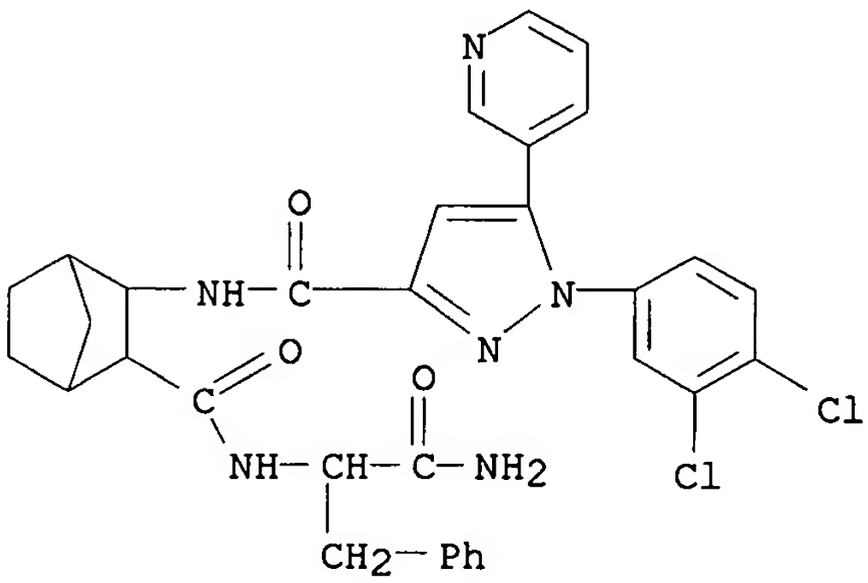
RN 627088-86-2 CAPLUS

CN Cyclohexanecarboxylic acid, [[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]- (9CI) (CA INDEX NAME)



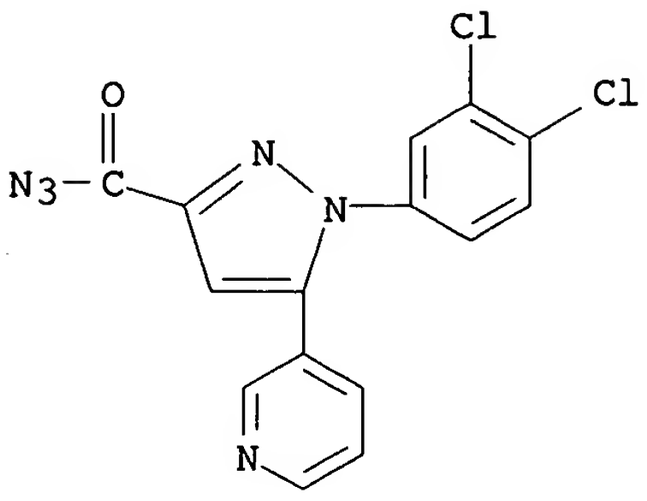
D1—CO_2H





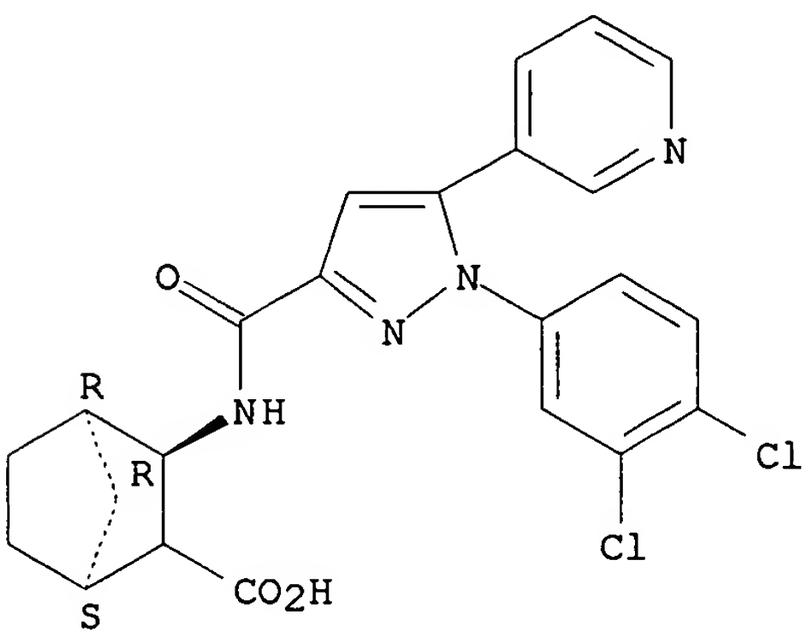
RN 627088-99-7 CAPLUS
CN 1H-Pyrazole-3-carboxamide, N-[3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)

IT **623158-60-1P 623158-63-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-60-1 CAPLUS
CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



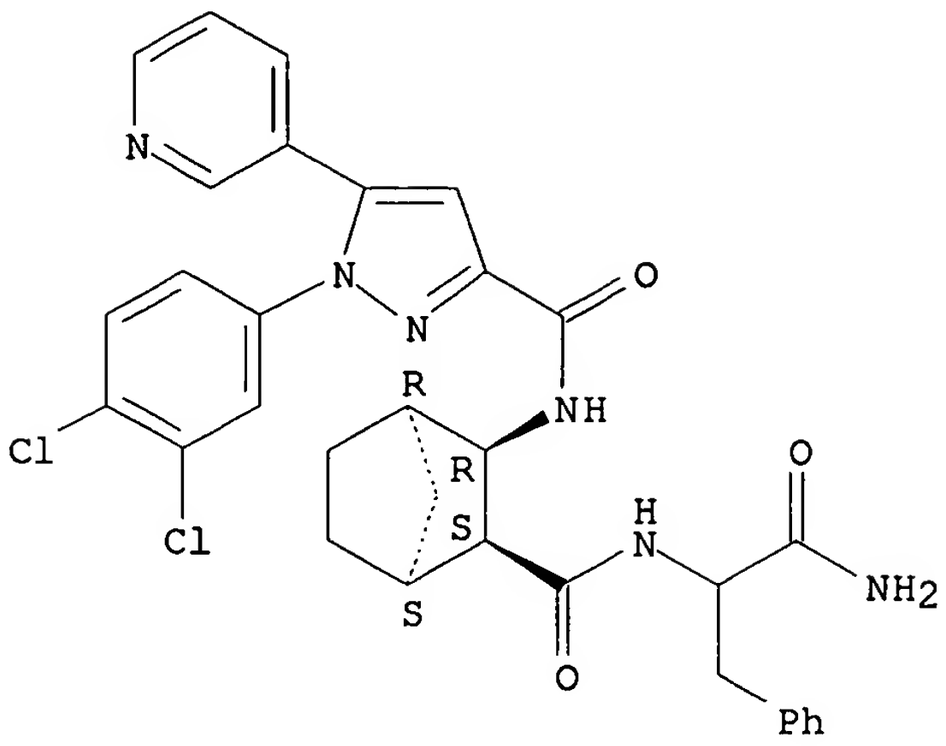
RN 623158-63-4 CAPLUS
CN Bicyclo[2.2.1]heptane-2-carboxylic acid, 3-[[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]-, (1R,3S,4S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



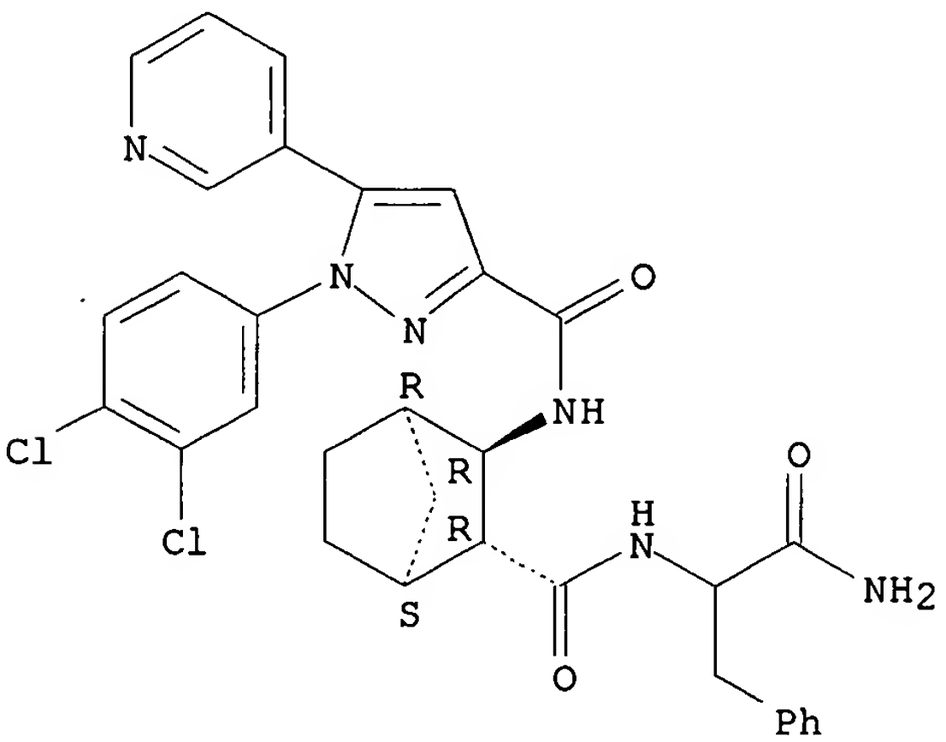
IT **623158-64-5P 623158-65-6P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-64-5 CAPLUS
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3S,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



RN 623158-65-6 CAPLUS
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3R,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:356451 CAPLUS
DOCUMENT NUMBER: 138:368907
TITLE: Preparation of pyrazolo[4,3-d]pyrimidin-7-ones as PDE9 inhibitors for treating cardiovascular disorders
INVENTOR(S): Deninno, Michael Paul; Hughes, Bernadette; Kemp, Mark Ian; Palmer, Michael John; Wood, Anthony
PATENT ASSIGNEE(S): Pfizer Limited, UK; Pfizer Inc.
SOURCE: PCT Int. Appl., 69 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003037899	A1	20030508	WO 2002-IB4385	20021022

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EP 1440073	A1	20040728	EP 2002-777623	20021022

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK

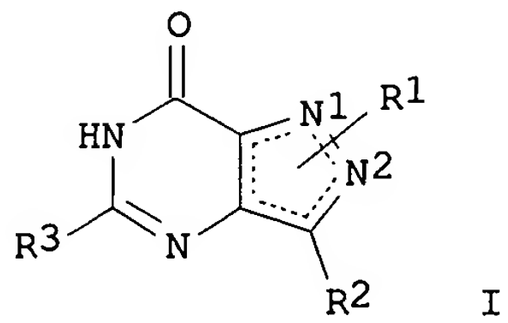
US 2003195205	A1	20031016	US 2002-283514	20021030

PRIORITY APPLN. INFO.:

GB 2001-26395	A	20011102
GB 2001-30695	A	20011221
GB 2002-16761	A	20020718
US 2002-350777P	P	20020122
US 2002-399905P	P	20020730
WO 2002-IB4385	W	20021022

OTHER SOURCE(S): MARPAT 138:368907

GI



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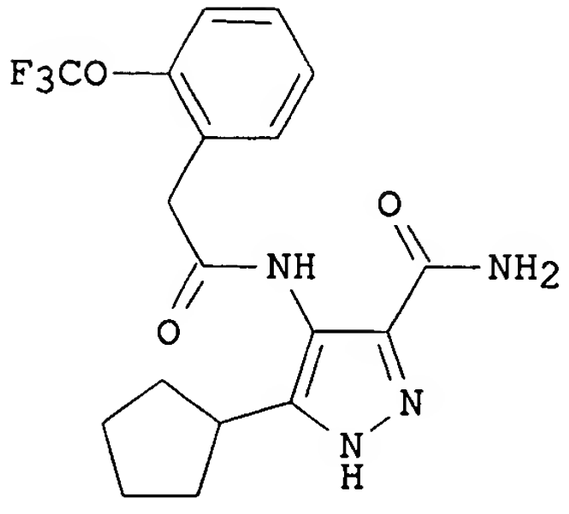
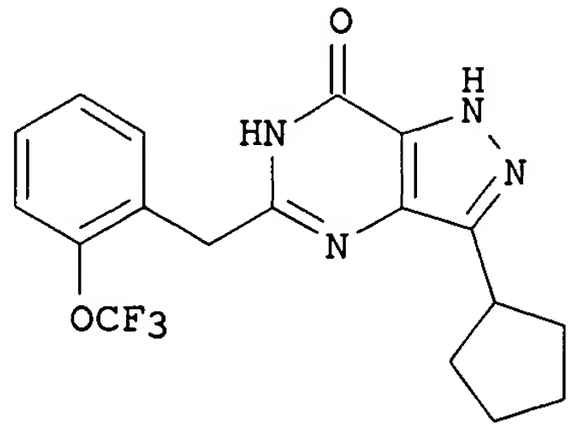


II



III

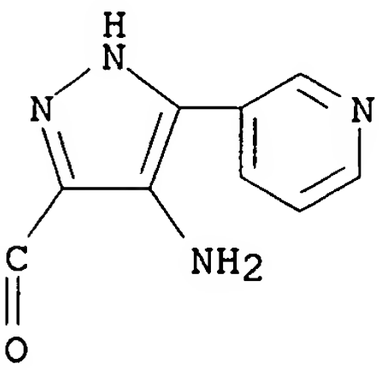
AB The title compds. [I; R1 = H, alkyl, wherein R1 is attached to either N1 or N2; R2 = alkyl, hydroxyalkyl, alkoxyalkyl, cycloalkyl, etc.; R3 = alkyl optionally substituted by (un)substituted Ph, cycloalkyl optionally substituted by alkyl, etc.], useful as PDE9 inhibitors for treating cardiovascular disorders, were prepd. and formulated. Thus, cyclization of the pyrazolecarboxamide II in the presence of tert-BuOK in iso-PrOH

afforded III which was found to have a greater than 40% inhibition against PDE9 at 1 .mu.M.

IT **265663-95-4P**, 4-Amino-5-(3-pyridyl)-1H-pyrazole-3-carboxamide
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of pyrazolo[4,3-d]pyrimidin-7-ones as PDE9 inhibitors for treating cardiovascular disorders)

RN 265663-95-4 CAPLUS

CN 1H-Pyrazole-3-carboxamide, 4-amino-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2003:356304 CAPLUS

DOCUMENT NUMBER: 138:368899

TITLE: Preparation of pyrazolopyrimidinones as PDE9 inhibitors for treatment of insulin resistance syndrome and type 2 diabetes

INVENTOR(S): Fryburg, David Albert; Gibbs, Earl Michael

PATENT ASSIGNEE(S): Pfizer Products Inc., USA

SOURCE: PCT Int. Appl., 104 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003037432	A1	20030508	WO 2002-IB3754	20020912

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

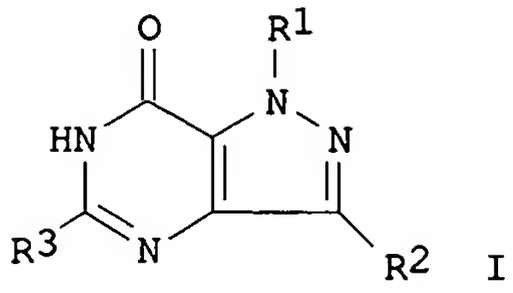
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004023989	A1	20040205	US 2002-283814	20021029

PRIORITY APPLN. INFO.: US 2001-336981P P 20011102

OTHER SOURCE(S): MARPAT 138:368899

GI



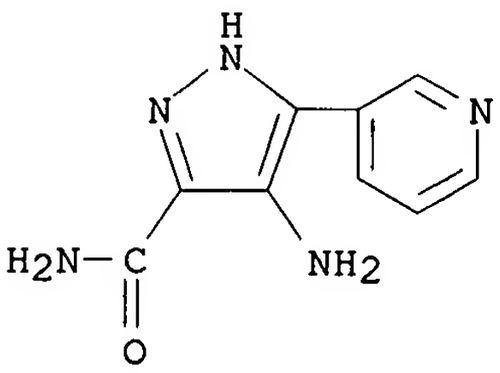
I

AB Pyrazolopyrimidinones I [R1 = H, alkyl; R2 = alkyl, cycloalkyl, heterocyclic; R3 = (un)substituted alkyl] were prepd. for use as PDE9 inhibitors in treating insulin resistance syndrome (IRS), hypertension and/or type 2 diabetes. Thus, Me2CHCOMe was treated with EtO2CCO2Et to give Me2CHCOCH2CO2Et which was cyclized with N2H4 to give Et 5-isopropyl-1H-pyrazole-3-carboxylate. This ester was hydrolyzed to the acid, nitrated, amidated, and reduced to give 4-amino-5-isopropyl-1H-pyrazole-3-carboxamide. Cyclization of this amide with 3-ClC6H4CH2CO2H gave I [R1 = H, R2 = CHMe2, R3 = 3-ClC6H4CH2] which reduced plasma glucose, triglycerides, and insulin at 10 mg/kg day for 5 days orally in mice.

IT **265663-95-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(prepn. of pyrazolopyrimidinones as PDE9 inhibitors for treatment of insulin resistance syndrome and type 2 diabetes)

RN 265663-95-4 CAPLUS

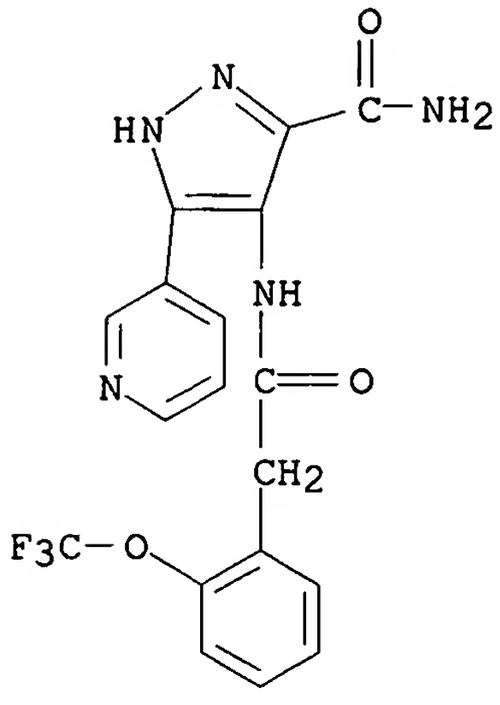
CN 1H-Pyrazole-3-carboxamide, 4-amino-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



IT **521300-42-5P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of pyrazolopyrimidinones as PDE9 inhibitors for treatment of insulin resistance syndrome and type 2 diabetes)

RN 521300-42-5 CAPLUS

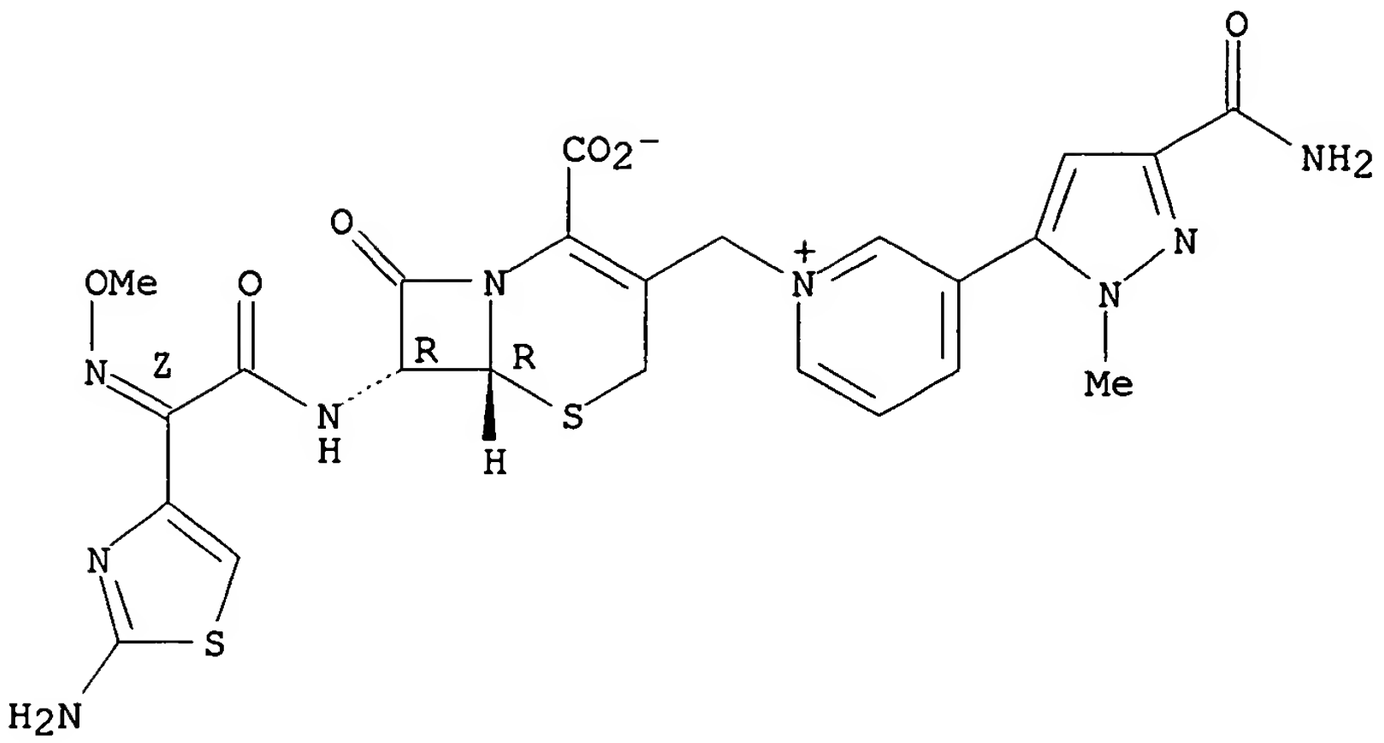
CN 1H-Pyrazole-3-carboxamide, 5-(3-pyridinyl)-4-[[[2-(trifluoromethoxy)phenyl]acetyl]amino]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

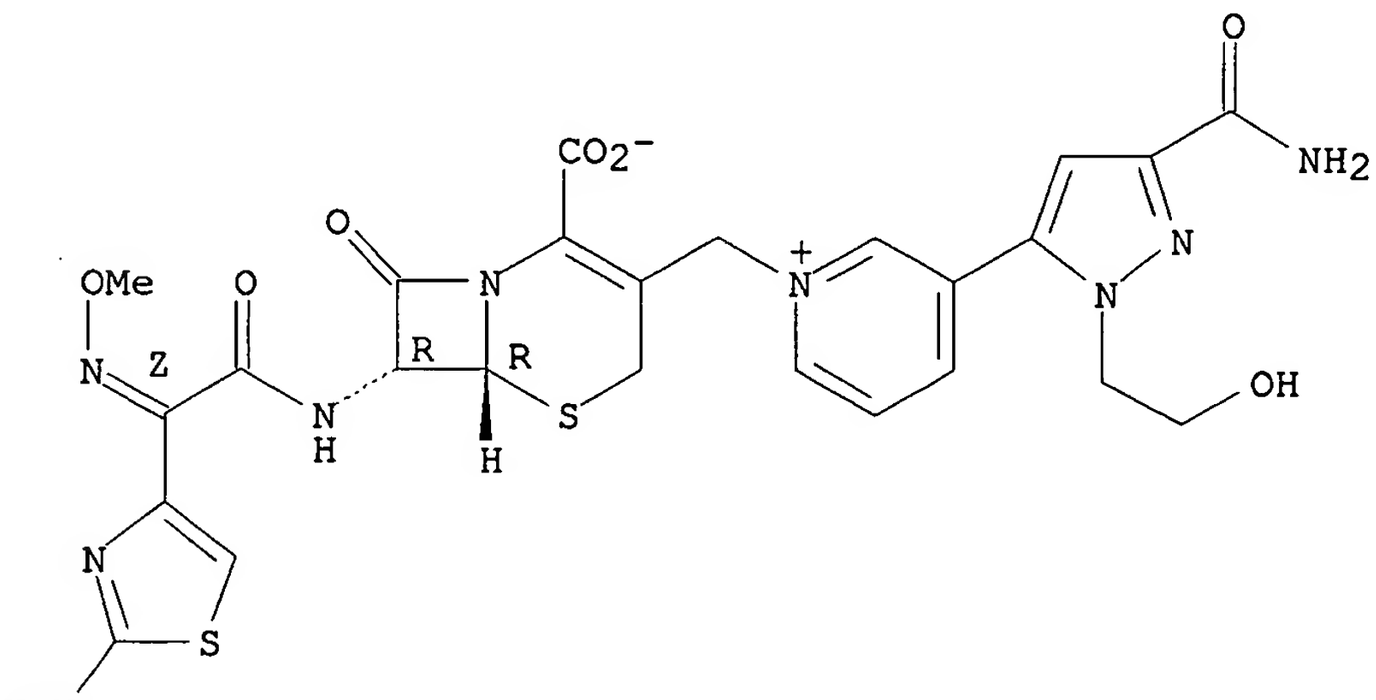
L4 ANSWER 8 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:419957 CAPLUS
DOCUMENT NUMBER: 133:150380
TITLE: Synthesis and structure-activity relationships of quaternary ammonium cephalosporins with 3-pyrazolylpyridinium derivatives
AUTHOR(S): Chang, Kwan Young; Kim, Sung Hoon; Nam, Ghilsoo; Seo, Jae Hong; Kim, Joong Hyup; Ha, Deok-Chan
CORPORATE SOURCE: Biochemicals Research Center, Korea Institute of Science and Technology, Seoul, 130-650, S. Korea
SOURCE: Bioorganic & Medicinal Chemistry Letters (2000), 10(11), 1211-1214
CODEN: BMCLE8; ISSN: 0960-894X
PUBLISHER: Elsevier Science Ltd.
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Cephalosporins with 3-pyrazolylpyridinium at C-3 position, which is supposed to exhibit synergic activity of ceftazidime and cefoselis, were synthesized and their antibacterial activity against Gram-pos. and Gram-neg. was inspected.
IT **287494-13-7P 287494-14-8P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(synthesis and structure-activity relationships of quaternary ammonium cephalosporins with 3-pyrazolylpyridinium derivs.)
RN 287494-13-7 CAPLUS
CN Pyridinium, 3-[3-(aminocarbonyl)-1-methyl-1H-pyrazol-5-yl]-1-[[(6R,7R)-7-[[(2Z)-(2-amino-4-thiazolyl)(methoxyimino)acetyl]amino]-2-carboxy-8-oxo-5-thia-1-azabicyclo[4.2.0]oct-2-en-3-yl]methyl]-, inner salt (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

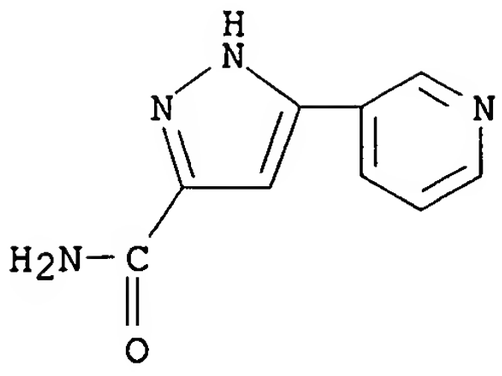


RN 287494-14-8 CAPLUS
CN Pyridinium, 3-[3-(aminocarbonyl)-1-(2-hydroxyethyl)-1H-pyrazol-5-yl]-1-[[(6R,7R)-7-[[(2Z)-(2-amino-4-thiazolyl)(methoxyimino)acetyl]amino]-2-carboxy-8-oxo-5-thia-1-azabicyclo[4.2.0]oct-2-en-3-yl]methyl]-, inner salt (9CI) (CA INDEX NAME)

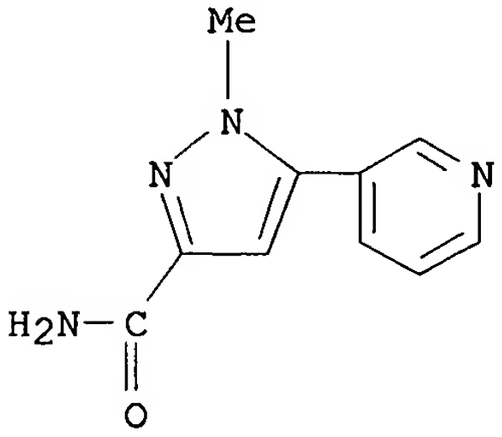
Absolute stereochemistry.
Double bond geometry as shown.



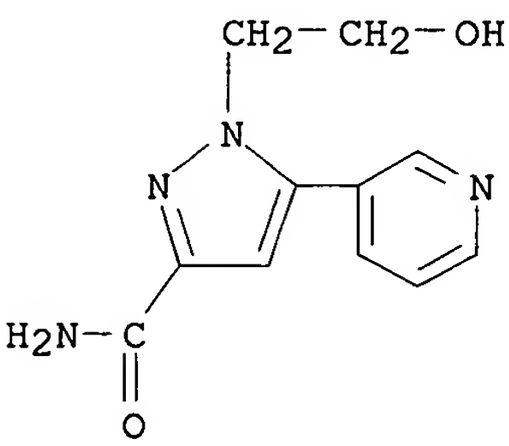
IT **287494-01-3P 287494-19-3P 287494-20-6P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis and structure-activity relationships of quaternary ammonium cephalosporins with 3-pyrazolylpyridinium derivs.)
RN 287494-01-3 CAPLUS
CN 1H-Pyrazole-3-carboxamide, 5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



RN 287494-19-3 CAPLUS
CN 1H-Pyrazole-3-carboxamide, 1-methyl-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



RN 287494-20-6 CAPLUS
CN 1H-Pyrazole-3-carboxamide, 1-(2-hydroxyethyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

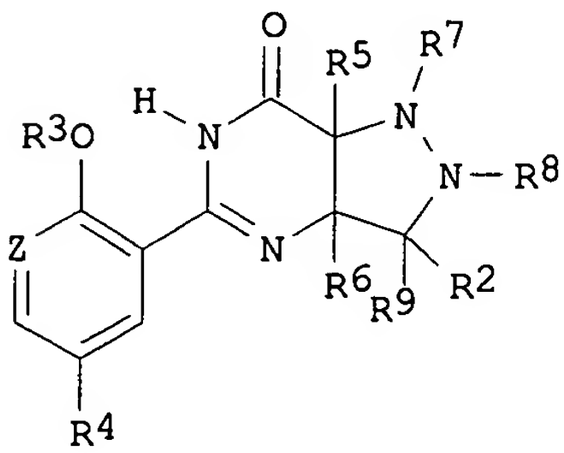
L4 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:291043 CAPLUS
DOCUMENT NUMBER: 132:308353
TITLE: Preparation of pyrazolopyrimidinones as cGMP phosphodiesterase inhibitors
INVENTOR(S): Bunnage, Mark Edward; Maw, Graham Nigel; Rawson, David James; Wood, Anthony; Mathias, John Paul; Street, Stephen Derek Albert
PATENT ASSIGNEE(S): Pfizer Limited, UK; Pfizer Inc.
SOURCE: PCT Int. Appl., 197 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

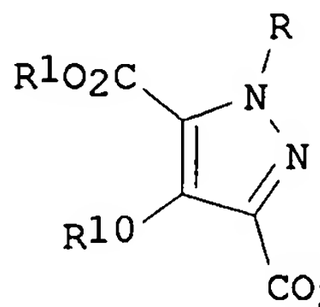
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000024745	A1	20000504	WO 1999-IB1706	19991019
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
AU 9959956	A1	20000515	AU 1999-59956	19991019
BR 9915532	A	20010814	BR 1999-15532	19991019
EP 1123296	A1	20010816	EP 1999-970992	19991019
EP 1123296	B1	20030917		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2002528456	T2	20020903	JP 2000-578315	19991019
AT 250063	E	20031015	AT 1999-970992	19991019
PT 1123296	T	20031231	PT 1999-970992	19991019
ES 2205945	T3	20040501	ES 1999-970992	19991019
US 6333330	B1	20011225	US 1999-426554	19991022

PRIORITY APPLN. INFO.:	GB 1998-23101	A	19981023	
	GB 1998-23102	A	19981023	
	WO 1999-IB1706	W	19991019	

OTHER SOURCE(S): MARPAT 132:308353
GI



I

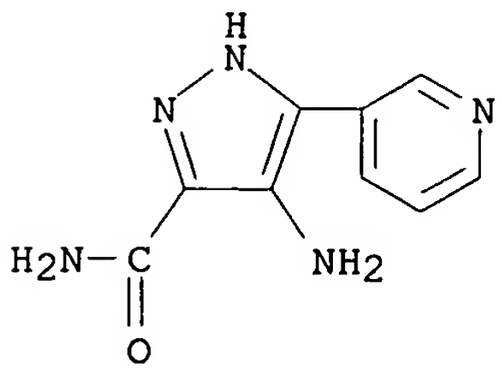


II

AB Title compds. [I; R2 = CONH2, CO2H, alkoxycarbonyl, (acyl)amino, etc.; R3 = H or (un)substituted alkyl; R4 = SO2NR14R15; R5R6 and R8R9 = bond and R7 = H, alkyl, heterocyclyl, aryl, etc.; R5R7 and R6R9 = bond and R8 = H, alkyl, heterocyclyl, aryl, etc.; NR14R15 = heterocyclyl; Z = CH or N] were prepd. for treatment of sexual dysfunction. Thus, pyrazole-3,5-dicarboxylic acid was nitrated and the product esterified to give pyrazolecarboxylate II (R = H, R1 = Me, R10 = NO2) which was N-alkylated by 2-chloromethylpyridine and the reduced product amidated by 2-(PrO)C6H4COCl to give II [R = 2-pyridylmethyl, R1 = Me, R10 = NHCOC6H4(OPr)-2]. The latter was heated with NH3 at 100.degree. to give I (R2 = CONH2, R3 = Pr, R5R6,R8R9 = bond, R7 = 2-pyridylmethyl)(III; R4 = H) which was converted to III (R4 = 4-methyl-1-pyrazinylsulfonyl). Data for biol. activity of I were given.

IT **265663-95-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)
(prepn. of pyrazolopyrimidinones as cGMP phosphodiesterase inhibitors)
RN 265663-95-4 CAPLUS
CN 1H-Pyrazole-3-carboxamide, 4-amino-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)

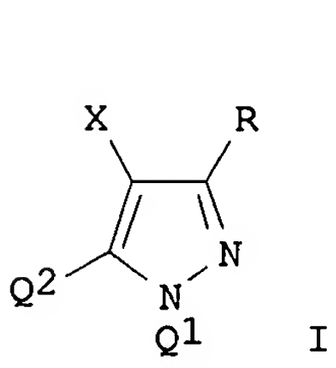


REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 10 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1996:281619 CAPLUS
DOCUMENT NUMBER: 124:317155
TITLE: Preparation of halopyrazolecarboxylic acids as herbicides
INVENTOR(S): Sato, Kazuo; Kudo, Noriaki; Pponma, Toyokuni; Endo, Takeshi; Kadotani, Junji; Horibe, Yoshimichi
PATENT ASSIGNEE(S): Sankyo Co, Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 26 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08012654	A2	19960116	JP 1994-144235	19940627

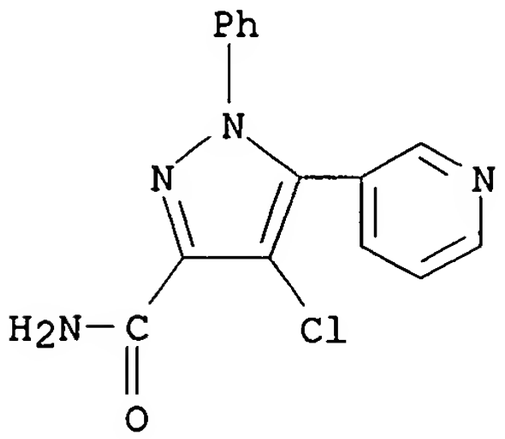
PRIORITY APPLN. INFO.: JP 1994-144235 19940627
OTHER SOURCE(S): MARPAT 124:317155
GI



I

AB The title compds. I [R = carboxyl, etc.; X = halo; Q1 = Ph, pyridinyl; Q2 = Ph, etc.] are prepd. I [X = Cl; Q1 = Q2 = phenyl; R = CO2Me] (m.p. 153 - 155.degree.) (at 10 g/are) gave 91 - 100% control of Echinochloa oryzicola and Scirpus juncoides and caused no damage to rice plants.
IT **176232-25-0P**
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of halopyrazolecarboxylic acids as herbicides)
RN 176232-25-0 CAPLUS

CN 1H-Pyrazole-3-carboxamide, 4-chloro-1-phenyl-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 11 OF 11 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1969:470595 CAPLUS
DOCUMENT NUMBER: 71:70595
TITLE: 5-Substituted pyrazole-3-carboxylic acid hydrazides
INVENTOR(S): Walker, Gordon Northrop
PATENT ASSIGNEE(S): CIBA Corp.
SOURCE: U.S., 3 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 3449350	A	19690610	US 1968-739135	19680624

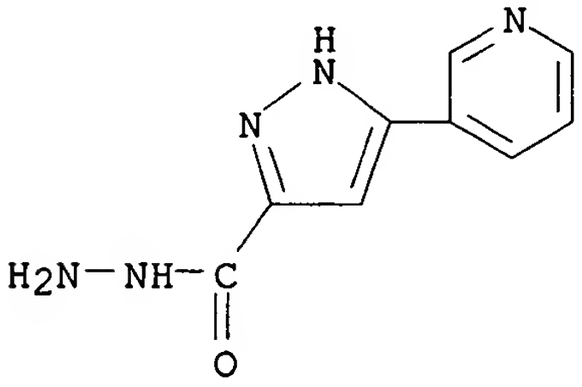
PRIORITY APPLN. INFO.: US 1968-739135 19680624
GI For diagram(s), see printed CA Issue.
AB Quaternary and addn. salts of the title compds. (I) with antitumor properties, are prepd. Thus, MeONa (from 5 g. Na) was suspended in 1 l. Et2O, 25.0 g. 1- acetylcyclohexene in 30.0 g. (CO2Et)2 added slowly and the mixt. kept at room temp. 2.5 days to give Et 2,3-dioxo-4-(1-cyclohexen-1-yl)butyrate (II). II (20.0 g.) and 200 cc. 5% aq. NaOH was stirred 15 min. at 90.degree., to give 2,3-dioxo-4-(1-cyclohex-1-enyl)butyric acid (III), m. 110-12.degree. (Et2O-petroleum ether). III (13.0 g.) in 100 cc. EtOH and 25 cc. 95% N2H4 was heated 30 min. at 95.degree. to give 5-(1-cyclohexen-1-yl)pyrazole-3-carboxylic acid (IV), m. 259-60.degree. (decompn.). IV (11.0 g.) and 120 cc. SOCl2 was refluxed 30 min. and evapd. in vacuo, 25 cc. 95% N2H4 added slowly and the mixt. heated at 5 min. 95.degree. to give I (R = 1-cyclohexen-1-yl) (Ia), m. 188-90.degree., also prepd. from 32.2 g. II and 150 cc. 95% N2H4 in 300 cc. EtOH heated 30 min. at 95.degree.. MeONa from 5.8 g. Na was added to 28.0 g. 3-acetylpyridine and 36.8 g. (CO2Et)2 in 50 cc. Et2O while stirring and cooling, the mixt. kept 3 days at room temp. and poured into 220 cc. ice water and the stirred soln. acidified with 18% aq. HCl to pH 5.5 to give Me 2,3-dioxo-3-(3-pyridyl)butyrate (V), m. 119-21.degree. (EtOH). Trans-esterification occurred during the reaction. A mixt. of 5.0 g. V, 1.6 g. NH2OH.HCl, and 50 cc. EtOH was refluxed 1 hr., to give 2-hydroxyimino-3-oxo-3-(3-pyridyl)butyrate (VI) hydrochloride, m. 201-2.degree. (EtOH-MeOH), 1.0 g. of which in the min. amt. EtOh was added to satd. aq. NaHCO3 to give the free base (VI), m. 133-5.degree. (MeOH-Et2O). VI (0.5 g.), 5 drops 95% N2H4, and 10 cc. EtOH was kept at room temp. overnight to give 1-hydroxyimino-3-oxo-3-(3-pyridyl)butyric acid hydrazide, m. 192-4.degree. (decompn.), 0.4 g. of which and 2 cc. 95%

N2H4 was heated at 95.degree. for 30 min. and evapd. in vacuo, the residue dissolved in 10 cc. MeOH contg. 0.5 cc. N2H4 and the soln. refluxed 1 hr. to give 5- (3-pyridyl)-pyrazole-3-carboxylic acid hydrazide (I, R = 3-pyridyl) (Ib), m. 263-5.degree. (decompn.) (MeOH). Ib was also prepd. from 11.6 g. V and 20 cc. 95% N2H4 heated in 200 cc. EtOH 10 min. at 95.degree..

IT **23424-35-3P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 23424-35-3 CAPLUS

CN Pyrazole-3-carboxylic acid, 5-(3-pyridyl)-, hydrazide (8CI) (CA INDEX NAME)



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CA INDEXING COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

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L1                  STR
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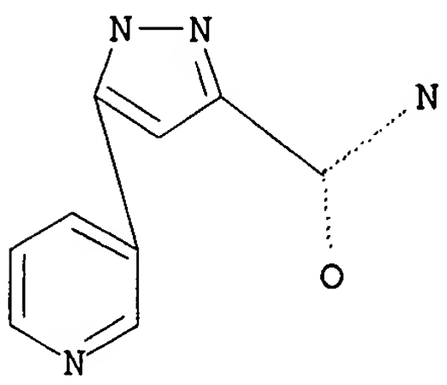

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L5             6 SEA L3


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L5   ANSWER 1 OF 6  USPATFULL on STN
ACCESSION NUMBER:         2004:152203  USPATFULL
TITLE:                    Prenylation inhibitors and methods of their synthesis
                          and use
INVENTOR(S):              Li, Francine Feirong, Raleigh, NC, UNITED STATES
                          Rehder, Kenneth S., Durham, NC, UNITED STATES
                          Campbell, Michael Gordon, Sagamore Hills, OH, UNITED
```

STATES
Viscardi, Celeste Patrice, Raleigh, NC, UNITED STATES
Strachan, Jon-Paul, Durham, NC, UNITED STATES
Guo, Zhengming, Raleigh, NC, UNITED STATES

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2004116425	A1	20040617	
APPLICATION INFO.:	US 2003-636327	A1	20030806	(10)

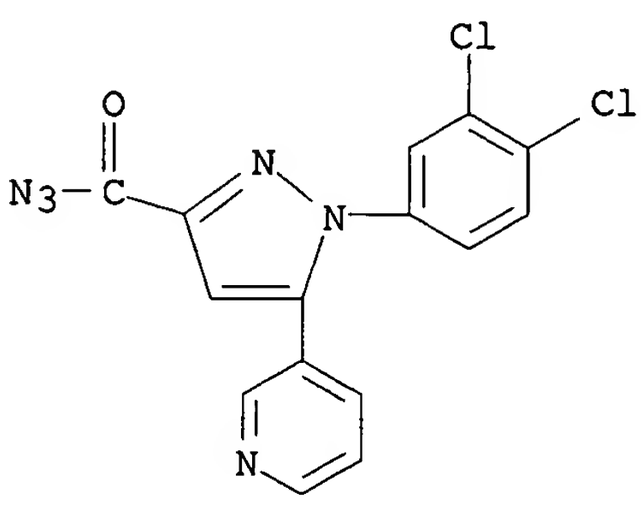
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2003-336285, filed on 3 Jan 2003, GRANTED, Pat. No. US 6649638
Continuation-in-part of Ser. No. US 2002-219628, filed on 14 Aug 2002, ABANDONED

	NUMBER	DATE
PRIORITY INFORMATION:	US 2003-454554P	20030314 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION
LEGAL REPRESENTATIVE: SHERIDAN ROSS PC, 1560 BROADWAY, SUITE 1200, DENVER, CO, 80202
NUMBER OF CLAIMS: 36
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 17 Drawing Page(s)
LINE COUNT: 2734
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention is directed to compounds useful in the treatment of diseases associated with prenylation of proteins and pharmaceutically acceptable salts thereof, to pharmaceutical compositions comprising same, and to methods for inhibiting protein prenylation in an organism using the same.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT **623158-60-1P**
(prepn. of substituted thiophenes and related compds. as prenylation inhibitors)
RN 623158-60-1 USPATFULL
CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



L5 ANSWER 2 OF 6 USPATFULL on STN
ACCESSION NUMBER: 2004:70742 USPATFULL
TITLE: Prenylation inhibitors and methods of their synthesis and use
INVENTOR(S): Brown, Bradley B., Durham, NC, UNITED STATES
Rehder, Kenneth S., Durham, NC, UNITED STATES

PATENT ASSIGNEE(S): PPD Discovery, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2004053970	A1	20040318
APPLICATION INFO.:	US 2003-646256	A1	20030822 (10)

RELATED APPLN. INFO.: Division of Ser. No. US 2003-336285, filed on 3 Jan 2003, GRANTED, Pat. No. US 6649638 Continuation-in-part of Ser. No. US 2002-219628, filed on 14 Aug 2002, ABANDONED
DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION
LEGAL REPRESENTATIVE: Gary J. Connell, SHERIDAN ROSS P.C., Suite 1200, 1560 Broadway, Denver, CO, 80202-5141
NUMBER OF CLAIMS: 62
EXEMPLARY CLAIM: 1
LINE COUNT: 1493

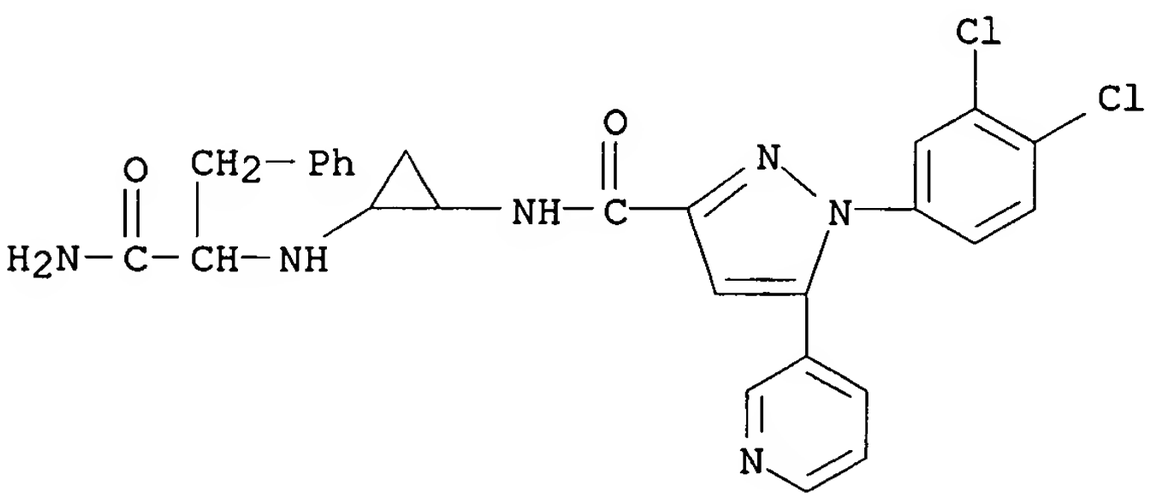
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention is directed to compounds useful in the treatment of diseases associated with prenylation of proteins and pharmaceutically acceptable salts thereof, to pharmaceutical compositions comprising same, and to methods for inhibiting protein prenylation in an organism using the same.

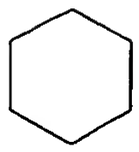
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT **623158-71-4P 627088-86-2P 627088-99-7P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)

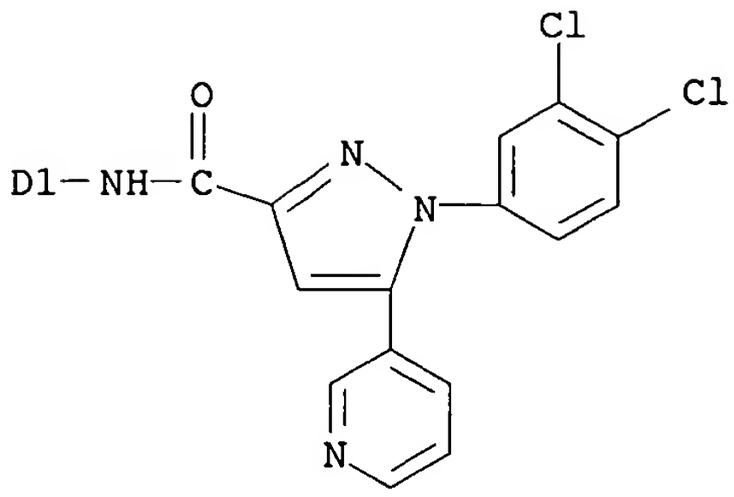
RN 623158-71-4 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[2-[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]cyclopropyl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



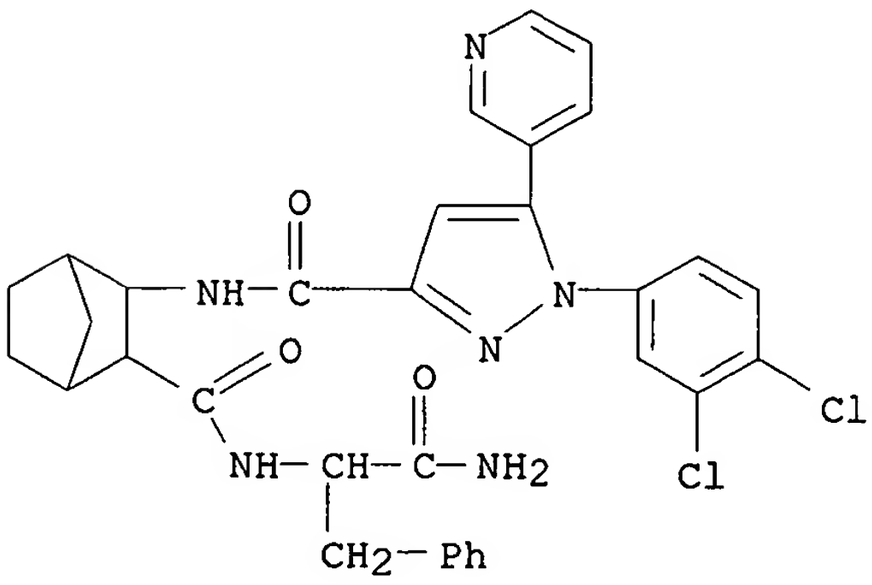
RN 627088-86-2 USPATFULL
CN Cyclohexanecarboxylic acid, [[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]- (9CI) (CA INDEX NAME)



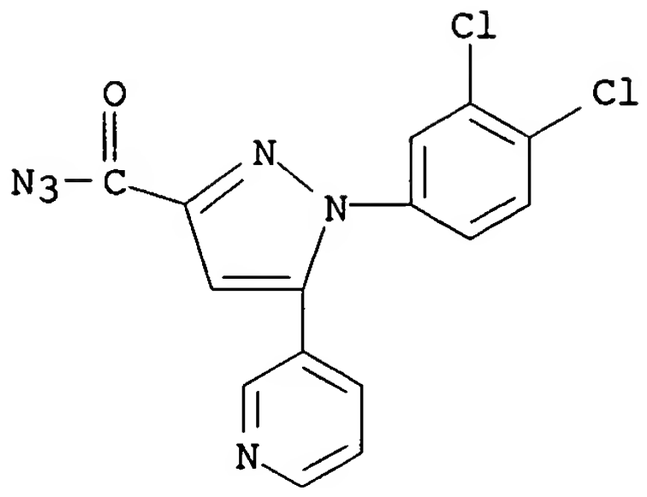
D1—CO_2H



RN 627088-99-7 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)

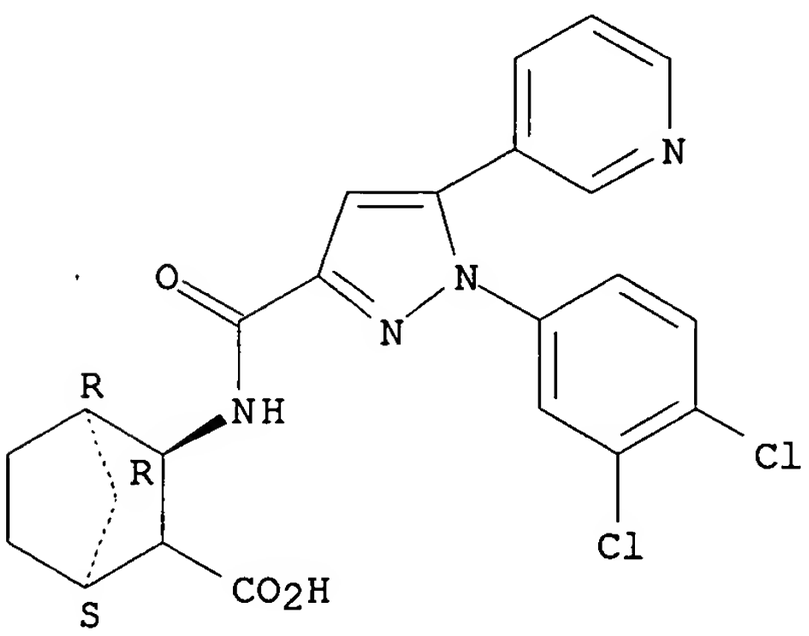


IT **623158-60-1P 623158-63-4P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-60-1 USPATFULL
CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



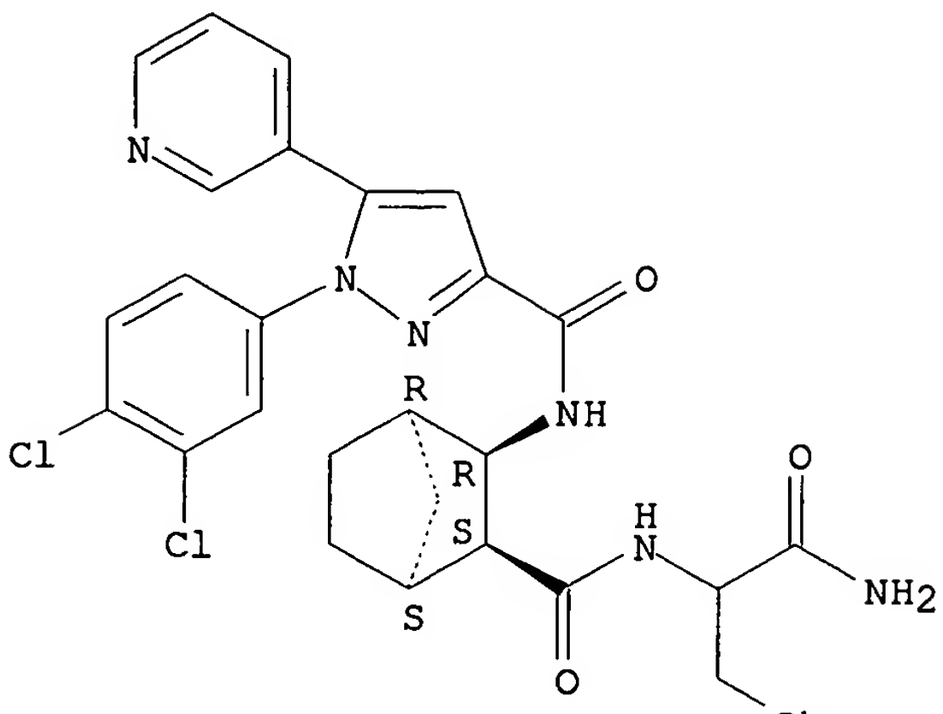
RN 623158-63-4 USPATFULL
CN Bicyclo[2.2.1]heptane-2-carboxylic acid, 3-[[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]-, (1R,3S,4S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



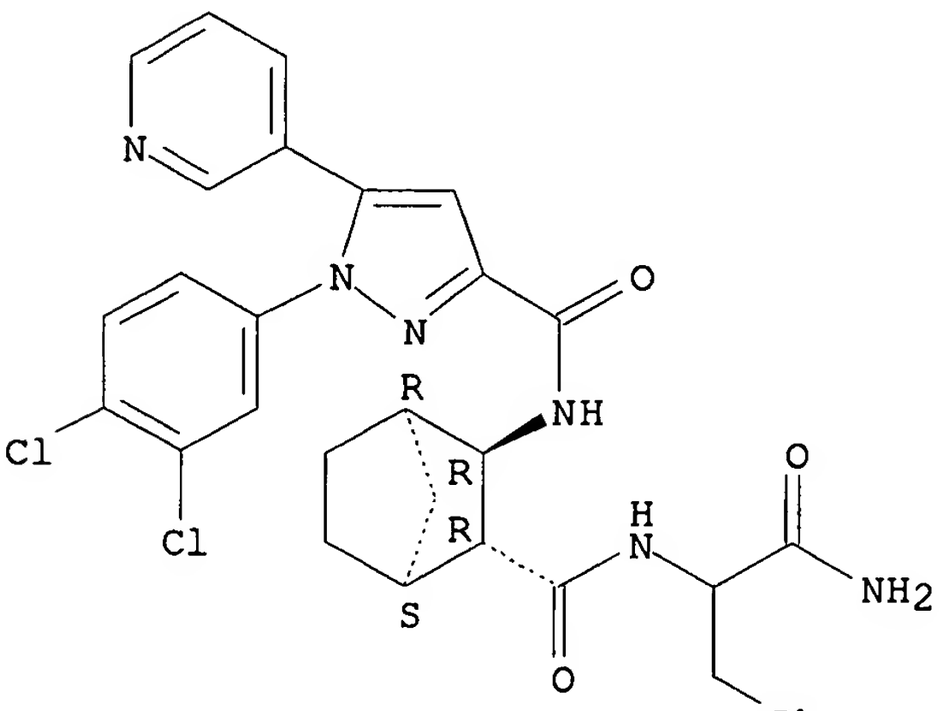
IT **623158-64-5P 623158-65-6P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-64-5 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3S,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



RN 623158-65-6 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3R,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



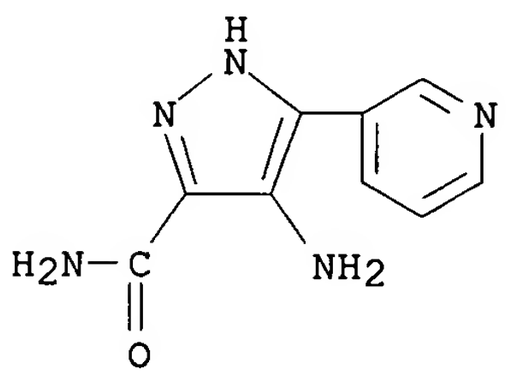
L5 ANSWER 3 OF 6 USPATFULL on STN

ACCESSION NUMBER:	2004:31847 USPATFULL			
TITLE:	Treatment of insulin resistance syndrome and type 2 diabetes with PDE9 inhibitors			
INVENTOR(S):	Fryburg, David A., East Lyme, CT, UNITED STATES Gibbs, Earl Michael, Oakdale, CT, UNITED STATES			
	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2004023989	A1	20040205	
APPLICATION INFO.:	US 2002-283814	A1	20021029	(10)
	NUMBER		DATE	
PRIORITY INFORMATION:	US 2001-336981P		20011102	(60)
DOCUMENT TYPE:	Utility			

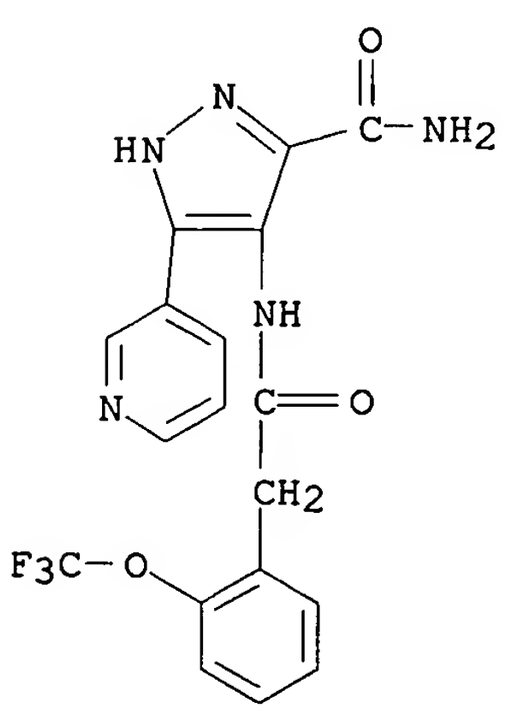
10/646,256

FILE SEGMENT: APPLICATION
LEGAL REPRESENTATIVE: PFIZER INC., PATENT DEPARTMENT, MS8260-1611, EASTERN POINT ROAD, GROTON, CT, 06340
NUMBER OF CLAIMS: 32
EXEMPLARY CLAIM: 1
LINE COUNT: 3315
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
AB This invention is directed to a method of treating insulin resistance syndrome (IRS), hypertension and/or type 2 diabetes in a mammal comprising administering to said mammal a cGMP PDE9 inhibitor or a pharmaceutical composition thereof. This invention is also directed to such methods wherein said cGMP PDE9 inhibitor is used in combination with other agents to treat IRS, hypertension and/or type 2 diabetes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT **265663-95-4**
(prepn. of pyrazolopyrimidinones as PDE9 inhibitors for treatment of insulin resistance syndrome and type 2 diabetes)
RN 265663-95-4 USPATFULL
CN 1H-Pyrazole-3-carboxamide, 4-amino-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



IT **521300-42-5P**
(prepn. of pyrazolopyrimidinones as PDE9 inhibitors for treatment of insulin resistance syndrome and type 2 diabetes)
RN 521300-42-5 USPATFULL
CN 1H-Pyrazole-3-carboxamide, 5-(3-pyridinyl)-4-[[[2-(trifluoromethoxy)phenyl]acetyl]amino]- (9CI) (CA INDEX NAME)



L5 ANSWER 4 OF 6 USPATFULL on STN
ACCESSION NUMBER: 2003:312771 USPATFULL
TITLE: Methods for treating carbonic anhydrase mediated

disorders

INVENTOR(S): Masferrer, Jaime L., Ballwin, MO, UNITED STATES
O'Neal, Janet M., St. Louis, MO, UNITED STATES
PATENT ASSIGNEE(S): Pharmacia Corporation (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003220376	A1	20031127
APPLICATION INFO.:	US 2003-367384	A1	20030214 (10)

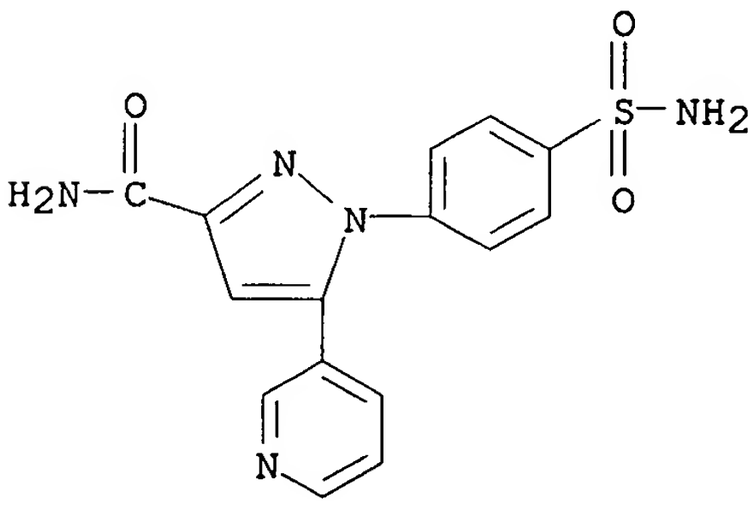
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2002-213793, filed on 7 Aug 2002, PENDING

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-311561P	20010810 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION
LEGAL REPRESENTATIVE: SENNIGER POWERS LEAVITT AND ROEDEL, ONE METROPOLITAN SQUARE, 16TH FLOOR, ST LOUIS, MO, 63102
NUMBER OF CLAIMS: 50
EXEMPLARY CLAIM: 1
LINE COUNT: 1946
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The current invention provides methods to treat or prevent carbonic anhydrase mediated diseases or disorders. The method generally comprises administering a tricyclic compound having a sulfonamide group to a subject wherein the compound inhibits carbonic anhydrase.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT **627094-49-9**
(sulfonamide-contg. cyclic compds. for treating carbonic anhydrase-mediated disorders, and use with other agents)
RN 627094-49-9 USPATFULL
CN 1H-Pyrazole-3-carboxamide, 1-[4-(aminosulfonyl)phenyl]-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



L5 ANSWER 5 OF 6 USPATFULL on STN
ACCESSION NUMBER: 2003:302881 USPATFULL
TITLE: Prenylation inhibitors and methods of their synthesis and use
INVENTOR(S): Brown, Bradley B., Durham, NC, United States
Rehder, Kenneth S., Durham, NC, United States
PATENT ASSIGNEE(S): PPD Discovery, Inc., Morrisville, NC, United States (U.S. corporation)

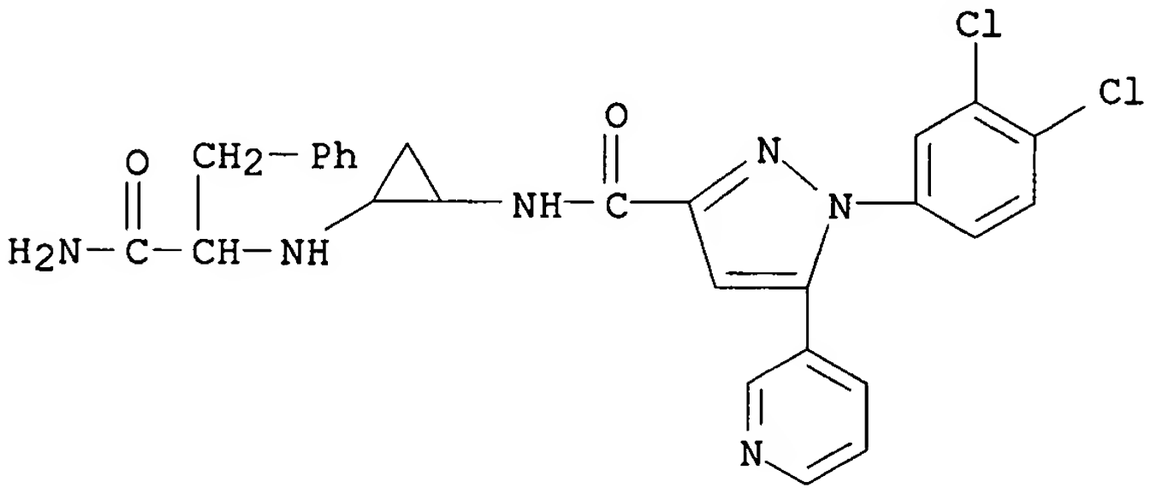
	NUMBER	KIND	DATE

PATENT INFORMATION:	US 6649638	B1	20031118
APPLICATION INFO.:	US 2003-336285		20030103 (10)

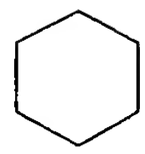
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 2002-219628, filed on 14 Aug 2002, now abandoned
DOCUMENT TYPE: Utility
FILE SEGMENT: GRANTED
PRIMARY EXAMINER: Fan, Jane
LEGAL REPRESENTATIVE: Sheridan Ross P.C.
NUMBER OF CLAIMS: 15
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)
LINE COUNT: 1348
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention is directed to compounds useful in the treatment of diseases associated with prenylation of proteins and pharmaceutically acceptable salts thereof, to pharmaceutical compositions comprising same, and to methods for inhibiting protein prenylation in an organism using the same.

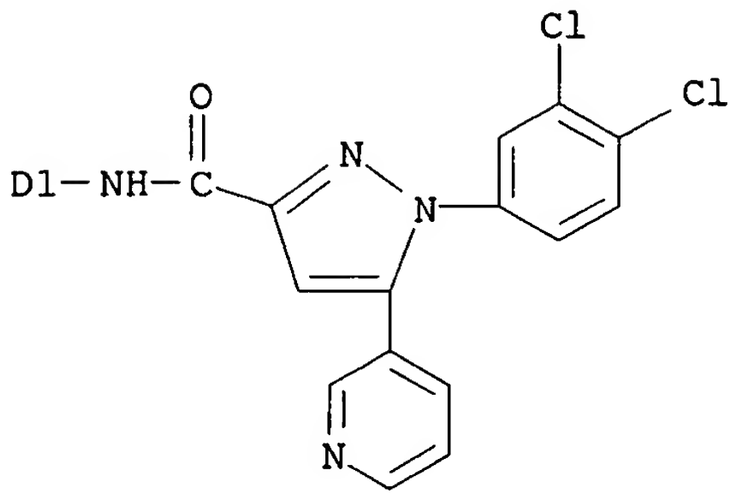
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT **623158-71-4P 627088-86-2P 627088-99-7P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-71-4 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[2-[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]cyclopropyl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



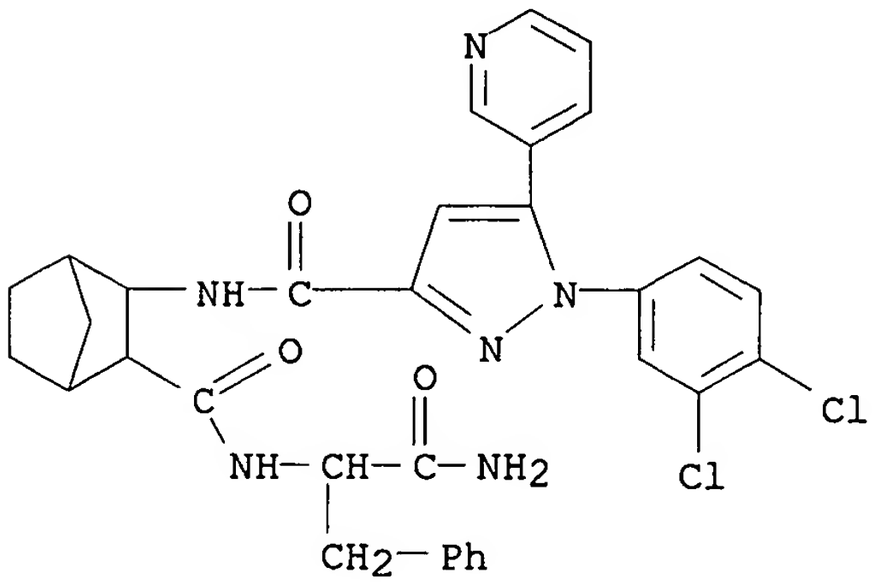
RN 627088-86-2 USPATFULL
CN Cyclohexanecarboxylic acid, [[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]- (9CI) (CA INDEX NAME)



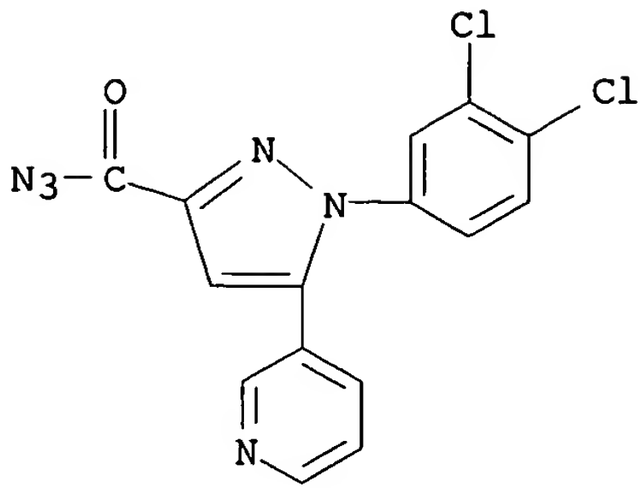
D1—CO_2H



RN 627088-99-7 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)

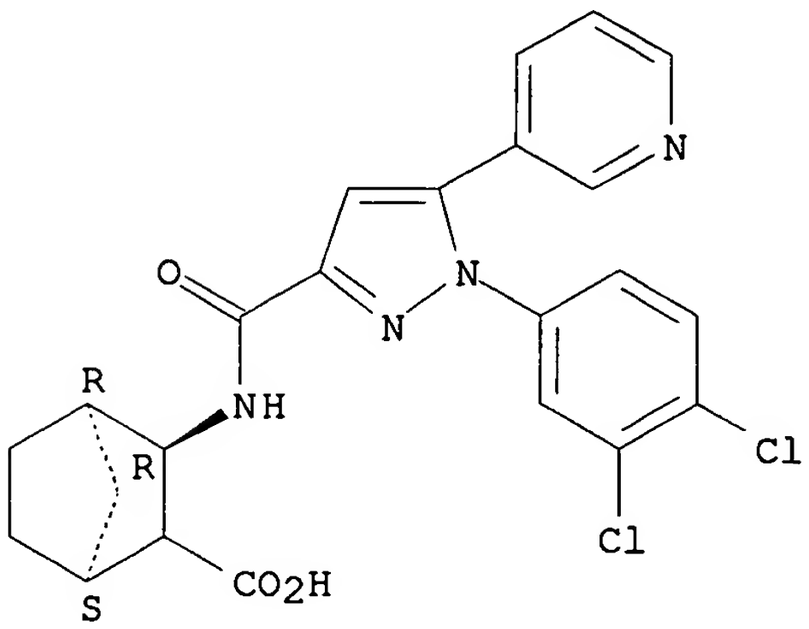


IT **623158-60-1P 623158-63-4P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-60-1 USPATFULL
CN 1H-Pyrazole-3-carbonyl azide, 1-(3,4-dichlorophenyl)-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)



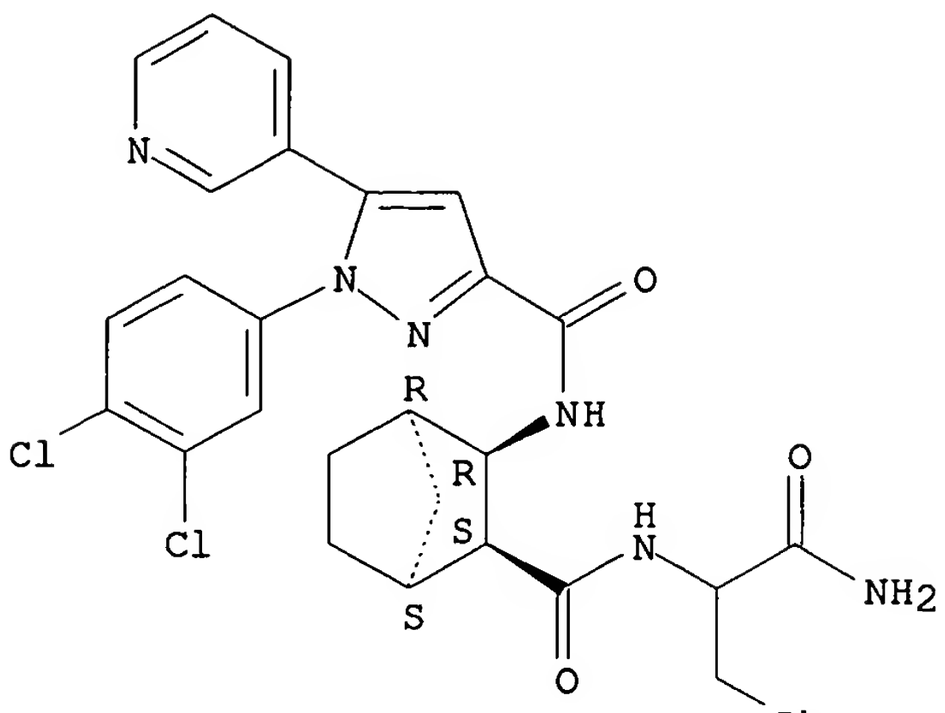
RN 623158-63-4 USPATFULL
CN Bicyclo[2.2.1]heptane-2-carboxylic acid, 3-[[[1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-1H-pyrazol-3-yl]carbonyl]amino]-, (1R,3S,4S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



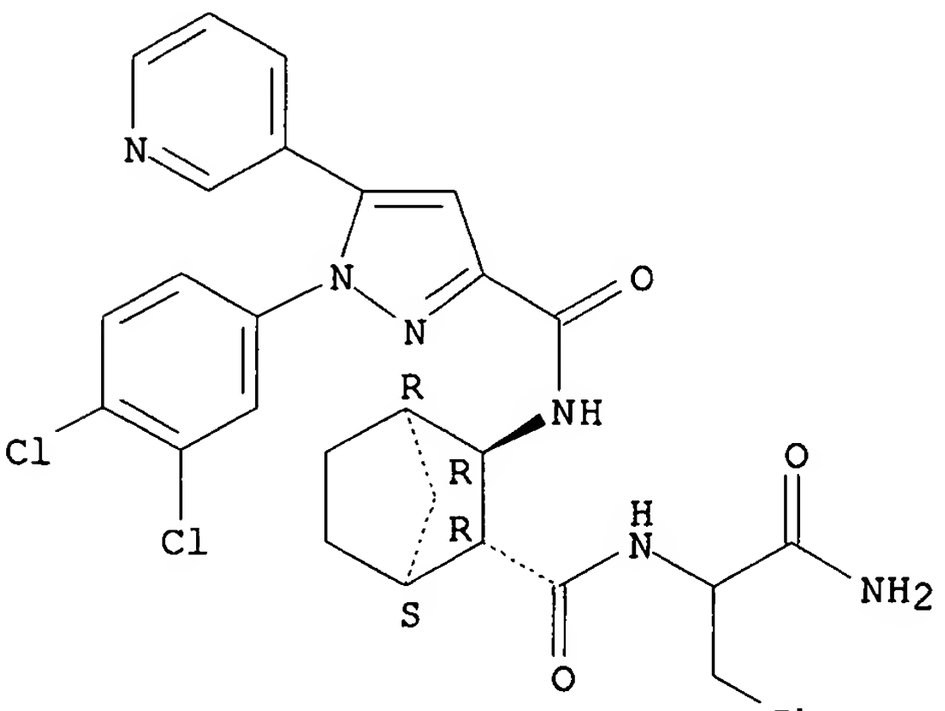
IT **623158-64-5P 623158-65-6P**
(prepn. of pyridylpyrazole peptide derivs. as prenylation inhibitors)
RN 623158-64-5 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3S,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



RN 623158-65-6 USPATFULL
CN 1H-Pyrazole-3-carboxamide, N-[(1R,2R,3R,4S)-3-[[[2-amino-2-oxo-1-(phenylmethyl)ethyl]amino]carbonyl]bicyclo[2.2.1]hept-2-yl]-1-(3,4-dichlorophenyl)-5-(3-pyridinyl)-, rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



L5 ANSWER 6 OF 6 USPATFULL on STN
ACCESSION NUMBER: 2003:277178 USPATFULL
TITLE: PDE9 inhibitors for treating cardiovascular disorders
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NUMBER OF CLAIMS:	26	
EXEMPLARY CLAIM:	1	
LINE COUNT:	1888	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to PDE9 inhibitors for treating cardiovascular disorders. Preferred PDE9 inhibitors are compounds of formula I wherein R.sup.1 is H or C.sub.1-6 alkyl, wherein R.sup.1 is attached to either N.sup.1 or N.sup.2; R.sup.2 is C.sub.1-6 alkyl optionally substituted by hydroxy or alkoxy; C.sub.3-7 cycloalkyl optionally substituted by alkyl, hydroxy or alkoxy; a saturated 5-6-membered heterocycle optionally substituted by alkyl, hydroxy or alkoxy; het1 or Ar.sup.1; R.sup.3 is C.sub.1-6 alkyl optionally substituted by 1 or 2 groups independently selected from: Ar.sup.2; C.sub.3-7cycloalkyl optionally substituted by C.sub.1-6alkyl; OAr.sup.2; SAr.sup.2; NHC(O)C.sub.1-6 alkyl; het.sup.2; xanthene; and naphthalene. ##STR1##

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IT **265663-95-4P**, 4-Amino-5-(3-pyridyl)-1H-pyrazole-3-carboxamide (prepn. of pyrazolo[4,3-d]pyrimidin-7-ones as PDE9 inhibitors for treating cardiovascular disorders)

RN 265663-95-4 USPATFULL

CN 1H-Pyrazole-3-carboxamide, 4-amino-5-(3-pyridinyl)- (9CI) (CA INDEX NAME)